
If both recessive alleles are present on respective homologous chromosomes, then the individual will be:

- A:
Heterozygous
- B:
Heterozygous recessive
- C:
Homozygous recessive
- D:
Homozygous dominant

Genetic make-up of a trait is called:

- A:
Phenotype
- B:
Genotype
- C:
Genome
- D:
Gene pool

In Mendel's monohybrid cross what percentage of round seed plants were produced by F₂ heterozygous round on self-fertilization?

- A :
25%
- B :
50%
- C :
75%
- D :
100%

A test cross is conducted to determine:

- A:
Genotype of a dominant phenotype
- B:
Genotype of a recessive phenotype
- C:
Phenotype of a dominant genotype
- D:
Phenotype of a recessive genotype

**If a trait is controlled by two or more than two genes,
then such genes are called as:**

- A:
Multiple alleles
- B:
Pleiotropic genes
- C:
Polygenes
- D:
Continuously varying traits

Which of the following is true about alleles?

- A :
They occupy different loci on same chromosome
- B :
They occupy same loci on different homologue
- C :
They occupy different loci on same homologue
- D :
They occupy same loci on respective homologue

An organism with two identical alleles for a trait is called:

- A:
Heterogenous
- B:
Homozygous
- C:
Co-dominant
- D:
Dominant

Seed shape in pea plant is:

A :
Trait

B :
Phenotype

C :
Genotype

D :
Genome

Dominance is physiological effect of an allele over its partner allele occupying:

- A:
Same locus on same chromosome
- B:
Same locus on respective homologue
- C:
Different locus on same chromosome
- D:
Different locus on respective homologue

_____ is the basic unit of biological information:

- A:
Gamete
- B:
Chromosome
- C:
DNA
- D:
Gene

Which of the following depicts the Mendel's dihybrid ratio?

A:
3 : 1

B:
9 : 3 : 3 : 1

C:
9 : 7

D:
15 : 1

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In dihybrid cross, out of 16 plants obtained, the number of genotypes will be:

- A :
4
- B :
9
- C :
16
- D :
12

_____ is a physiological effect of an allele over its partner allele on same gene locus.

- A :
Epistasis
- B :
Dominance
- C :
Bombay phenotype
- D :
Gene linkage

Eye colour in heterozygous *Drosophila* with genotype w^+w is an example of:

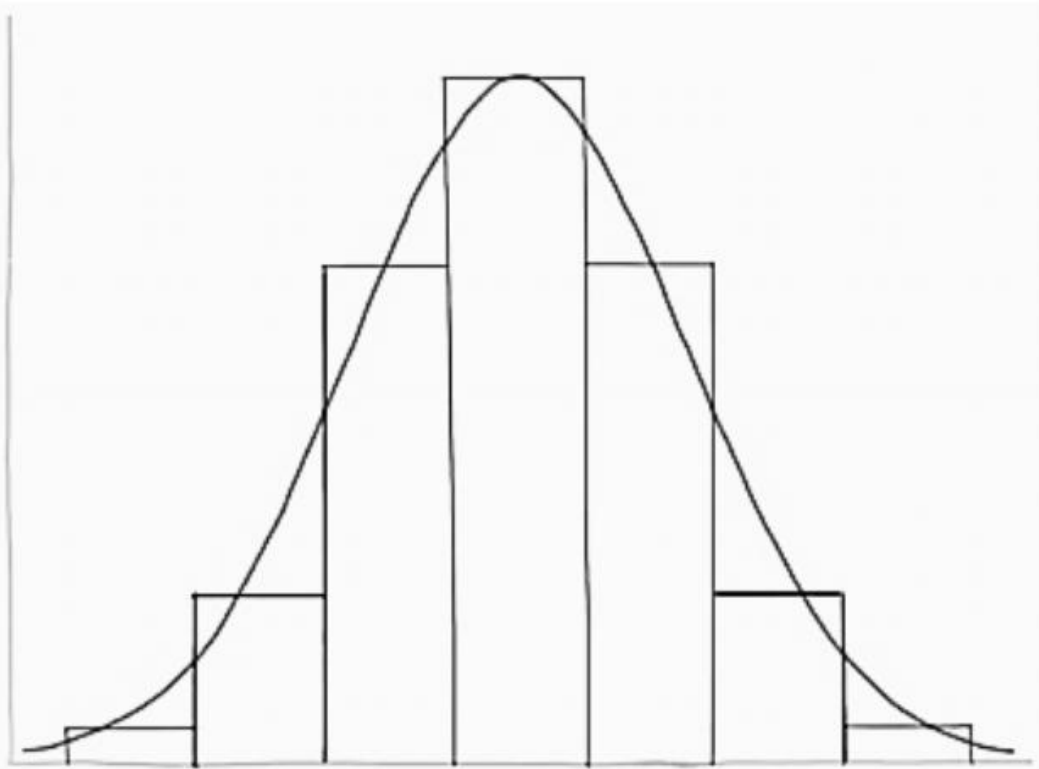
- A:
Complete dominance
- B:
Incomplete dominance
- C:
Co-dominance
- D:
Over dominance

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Intelligence is an example of:

- A:
Dominance
- B:
Pleiotropy
- C:
Epistasis
- D:
Polygenic inheritance

Which of the following trait can be related with given graph



- A:
Tongue rolling
- B:
ABO Blood Group
- C:
Seed colour in pea plant
- D:
Eye colour in human

Multiple alleles are the alleles of a gene and are always:

- A:
More than 1
- B:
More than 2
- C:
More than 3
- D:
More than 4

Types of blood group A, B, O and AB are known as:

- A:
Phenotypes
- B:
Genomes
- C:
Genotypes
- D:
Multiple alleles

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Which of the following blood group in humans is an example of co-dominance?

A :
A

B :
AB

C :
B

D :
O

Example of autosomal recessive trait:

- A:
Haemophilia A
- B:
Haemophilia B
- C:
Blue blindness
- D:
Hypophosphatemia

Which one is mismatched?

- A:
Haemophilia C – autosomal
- B:
Blue opsin - autosome 7
- C:
tfm – X chromosome
- D:
Pattern baldness – X linked

Partially functional opsins are present in

- A:
Protanopia
- B:
Deutanopia
- C:
Red green colorblindness
- D:
Protanomalous

Most prevalent abnormality of blood clotting factor is of:

A:
Factor VII

B:
Factor VIII

C:
Factor IX

D:
Factor X

Chromosomal combination of a person with tfm syndrome is:

A:
XO

B:
XY

C:
XX

D:
XYY

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_____ gene is male sex switch which triggers developmental process towards maleness:

A:
Se

B:
SRY

C:
tfm

D:
H

Linked genes can be separated through:

- A:
Segregation of alleles
- B:
Independent assortment
- C:
Crossing over
- D:
Mutation

Gene for formation of blue opsins is located on:

- A:
Chromosomes # 19

- B:
Chromosome # 9

- C:
Chromosome # 7

- D:
Chromosome # 11

It represents normal colour vision:

A :
Monochromacy

B :
Dichromacy

C :
Trichromacy

D :
Tetrachromacy

Genes for which of the following form a linkage group on chromosome 11?

- A:
Leukemia, Albinism
- B:
Hemophilia, Gout
- C:
Gout, Sickle cell anemia
- D:
Color blindness, Hemophilia

Hemophilia B is due to absence or abnormality of blood clotting factor:

A:
VIII

B:
IX

C:
X

D:
XI

A dichromate is unable to perceive:

- A:
One primary colour
- B:
Three primary colours
- C:
Two primary colours
- D:
All the colours

Women with normal colour vision whose father was red-green colour blind married a red-green colour blind man. What is the probability of her first born child being red-green colourblind:

- A:
1.0
- B:
0.75
- C:
0.66
- D:
0.50

Two normal parents have an albino child. What is the probability that their next child will also be an albino?

- A :
0%
- B :
50%
- C :
25%
- D :
100%

A boy receives his X-chromosome from:

- A:
His mother only
- B:
Both father and mother
- C:
His father only
- D:
Either father or mother

A hereditary disease which is never passed on from father to son is:

- A :
Y- linked disease
- B :
Autosomal disease
- C :
X- linked disease
- D :
None of these

Haemophilia is caused by a sex-linked, recessive allele. Two parents have a haemophiliac son, a normal son and a haemophiliac daughter.

What are the most likely genotypes of the parents?

- A:
- | | |
|-----------|---------|
| Mother | Father |
| $X^H X^h$ | $X^h Y$ |
- B:
- | | |
|-----------|---------|
| Mother | Father |
| $X^H Y^h$ | $X^H Y$ |
- C:
- | | |
|-----------|---------|
| Mother | Father |
| $X^h X^h$ | $X^H Y$ |
- D:
- | | |
|---------|---------|
| Mother | Father |
| $X^H Y$ | $X^h Y$ |

A normal woman whose father was red-blind marries a red-blind man. What proportion of their children can have normal colour vision?

- A :
25%
- B :
75%
- C :
50%
- D :
100%

A person having recessive alleles for blue opsins on autosome 7 can perceive:

- A:
Blue color
- B:
Red and green colors
- C:
All three primary colors
- D:
Red and blue colors

Which of the following is X-linked dominant trait in humans?

- A:
Hemophilia A
- B:
Vitamin D resistant rickets
- C:
Red-green colour blindness
- D:
Testicular feminization syndrome

Haemophilia can be the result of:

- A :
Reduction of blood clotting factors
- B :
Complete absence of blood clotting factors
- C :
Malfunctioning of blood clotting factors
- D :
All A, B, C

If a carrier woman for haemophilia is married to a normal man, then all of the following combinations can exist except:

A:
 $X^H X^H$

B:
 $X^H Y$

C:
 $X^h Y$

D:
 $X^h X^h$

If a female is colour blind having genetic combination X^cX^c , then the possible combination of her parents should be:

A:
 X^CX^C, X^CY

B:
 X^CX^C, X^cY

C:
 X^cX^c, X^cY

D:
 X^CX^c, X^CY

This is not true about testicular feminization syndrome:

- A:
It is androgen insensitivity syndrome
- B:
Individuals are females in appearance
- C:
Having blind vagina but no uterus
- D:
Degenerated testes are present in scrotum

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A person can differentiate between colours due to presence of normal:

- A:
Rod cells in retina
- B:
Cone cells in retina
- C:
Rod cells in cornea
- D:
Cone cells in cornea

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It passes directly from father to son:

- A:
X-linked trait
- B:
Y-linked trait
- C:
X-linked recessive trait
- D:
X-linked dominant trait

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Gene for blood clotting factor XI is located on:

- A:
X chromosome
- B:
Y chromosome
- C:
Autosome
- D:
Both X & Y chromosomes

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Physical association of two genes is known as:

- A:
Heterozygous
- B:
Recombination
- C:
Linkage
- D:
Dominance

If a carrier woman for haemophilia is married to a normal man, then all of the following combinations can exist in progeny except:

A:
 $X^H X^H$

B:
 $X^H Y$

C:
 $X^h Y$

D:
 $X^h X^h$

Genes present on both X and Y chromosomes are known as:

- A:
X-linked
- B:
Y-linked
- C:
Sex linked
- D:
Pseudoautosomal

Even a single recessive allele on X chromosome in male can be expressed because:

- A :
All genes on X chromosome can be expressed
- B :
Y chromosome has dominant allele for that trait
- C :
X chromosome alleles are dominant over Y chromosome
- D :
Y chromosome does not have counterpart of allele of X chromosome